In the Claims:

Please cancel claims 1-11, 14, 17-19 and 22-35 without prejudice thereto. Please amend the claims as follows:

Claims 1 - 11 (canceled)

- 12. (currently amended) A method for regulating of increasing I-BABP expression in a mammal which comprises activating or inhibiting the Farnasoid X Receptor comprising administering an effective amount of a non-steroidal Farnesoid X Receptor agonist to said mammal.
- 13. (currently amended) The method of claim 12, which comprises binding an activating amount of chenodeoxycholic acid to the Farnesoid X Receptor where said Farnesoid X Receptor agonist is GW4064.
 - 14. (canceled)
- 15. (currently amended) A method of regulating the bile transport system increasing bile acid transport from the intestinal lumen across the intestinal membrane in a mammal which comprises activating the Farnesoid X Receptor with a binding ligand administering an effective amount of a non-steroidal Farnesoid X Receptor agonist to said mammal.
- 16. (currently amended) The method of claim 15, wherein said FXR agonist is GW4064 the binding ligand is GW4064, chenodeoxycholic acid, lithocholic acid, deoxycholic acid, or a glycine or taurine conjugated conjugate derivative thereof.
 - 17-19 (cancelled)
- 20. (currently amended) A method of <u>lowering serum triglycerides</u> treating in a mammal a disease which is affected by cholesterol, triglyceride, or bile acid levels

comprising administering to a mammal in need of such treatment a therapeutically effective an effective serum triglyceride lowering amount of a non-steroidal agonist ligand for Farnesoid X Receptor.

21. (currently amended) A method of treating-atherosclerosis, gallstone disease, lipid disorders, obesity or a cardiovascular disorder in a mammal in need of such treatment, comprising administration of a therapeutically effective amount of a compound which was identified by the method of Claim 4 non-steroidal FXR agonist.

22-35 (canceled)